Hess's Law

Advanced Chemistry

Introduction

It is often possible to calculate ΔH for a reaction from the tabulated ΔH values of other reactions.

Thus, it is not necessary to make calorimetric measurements for all reactions.

Enthalpies in Reactions

- Because enthalpy is a state function, the enthalpy change, ΔH, associated with any chemical process depends only on the amount of matter that undergoes change and on the nature of the initial state of reactants and final state of products.
 - Whether a reaction is carried out in one step or in a series of steps, the sum of the enthalpy changes associated with individual steps must be the same as the enthalpy change associated with a one-step process.

Hess's Law

Hess's Law states that if a reaction is carried out in a series of steps, ΔH for the overall reaction equals the sum of the enthalpy changes for the individual steps.

3 Guidelines When Using Thermochemical Equations

1. Enthalpy is an extensive property.

The magnitude of ΔH is proportional to the amount of reactant consumed in the process.

Example: 890 kJ of heat is produced when 1 mole of CH_4 is burned in a constant-pressure system:

 $CH_4(g) + 2O_2(g) \rightarrow CO_2(g) + 2H_2O(l) \quad \Delta H = -890kJ$

▶ Because the combustion of 1mol of CH_4 with 2 mol of O_2 releases 890 kJ of heat, the combustion of 2 moles of CH_4 and 4 moles of O_2 releases twice as much heat, 1780 kJ

3 Guidelines When Using Thermochemical Equations

2. The enthalpy change for a reaction is equal in magnitude, but opposite to sign to delta H for the reverse reaction.

CH₄(g) + 2O₂(g) → CO₂(g) + 2H₂O(l) Δ H = -890kJ CO₂(g) + 2H₂O(l) → CH₄(g) + 2O₂(g) Δ H = +809 kJ

3 Guidelines When Using Thermochemical Equations

3. The enthalpy change for a reaction depends on the states of the reactants and products

- If the product from equation were H₂O(g) instead of H₂O(l), ΔHrxn would be -802 kJ instead of -890 kJ.
- Less heat would be available for transfer to the surroundings because the enthalpy of $H_2O(g)$ is greater than that of $H_2O(l)$

In other words

 $2H_2O(l) \rightarrow 2H_2O(g) \quad \Delta H = +88kJ$

Example

 $\begin{array}{ll} \mathsf{CH}_4(\mathsf{g}) + 2\mathsf{O}_2(\mathsf{g}) \rightarrow \mathsf{CO}_2(\mathsf{g}) + 2\mathsf{H}_2\mathsf{O}(\mathsf{g}) & \Delta\mathsf{H} = -802 \text{ kJ} \\ \bullet & 2\mathsf{H}_2\mathsf{O}(\mathsf{g}) \rightarrow 2\mathsf{H}_2\mathsf{O}(\mathsf{l}) & \Delta\mathsf{H} = -88 \text{ kJ} \\ \hline \mathsf{CH}_4(\mathsf{g}) + 2\mathsf{O}_2(\mathsf{g}) \rightarrow \mathsf{CO}_2(\mathsf{g}) + 2\mathsf{H}_2\mathsf{O}(\mathsf{l}) & \Delta\mathsf{H} = -890 \text{ kJ} \end{array}$

*Compounds on opposite sides of arrows are subtracted, while objects on same side are added.

Example

The enthalpy of reaction for the combustion of C to CO₂ is -393.5 kJ/mol C, and the enthalpy for the combustion of CO to CO₂ is -283.0 kJ/mol CO:

 $C(s) + O_2(g) \rightarrow CO_2(g) \quad \Delta H = -393.5 \text{ kJ}$

■ $CO(g) + \frac{1}{2} O_2(g) \rightarrow CO_2(g)$ $\Delta H = -283.0 \text{ kJ}$

 $C(s) + \frac{1}{2} O_2(g) \rightarrow CO(g) \qquad \Delta H = ?$

More Practice

Carbon occurs in two forms, graphite and diamond. The enthalpy of the combustion of graphite is -393.5 kJ/mol, and that of diamond is -395.4 kJ/mol:

C(graphite) + $O_2(g) \rightarrow CO_2(g)$ $\Delta H = -393.5 \text{ kJ}$ ← C(diamond) + $O_2(g) \rightarrow CO_2(g)$ $\Delta H = -395.4 \text{ kJ}$

C(graphite) \rightarrow C(diamond) Δ H = ?

More Practice $\Delta H = -1299.6 \text{ kJ}$ $C_2H_2(g) + 5/2 O_2(g) \rightarrow 2CO_2(g) + H_2O(l)$ $\Delta H = -393.5 \, kJ$ $C(s) + O_2(g) \rightarrow CO_2(g)$ $\Delta H = -285.8 \, kJ$ $2C(s) + H_2(g) \rightarrow C_2H_2(g)$ $\Delta H = ?$

Practice

$NO(g) + O_3(g) \rightarrow NO_2(g) + O_2(g) \qquad \Delta H = -198.9 \text{ kJ}$ $O_3(g) \rightarrow 3/2 O_2(g) \qquad \Delta H = -142.3 \text{ kJ}$ $O_2(g) \rightarrow 2O(g) \qquad \Delta H = 495.0 \text{ kJ}$ $NO(g) + O(g) \rightarrow NO_2(g) \qquad \Delta H = ?$